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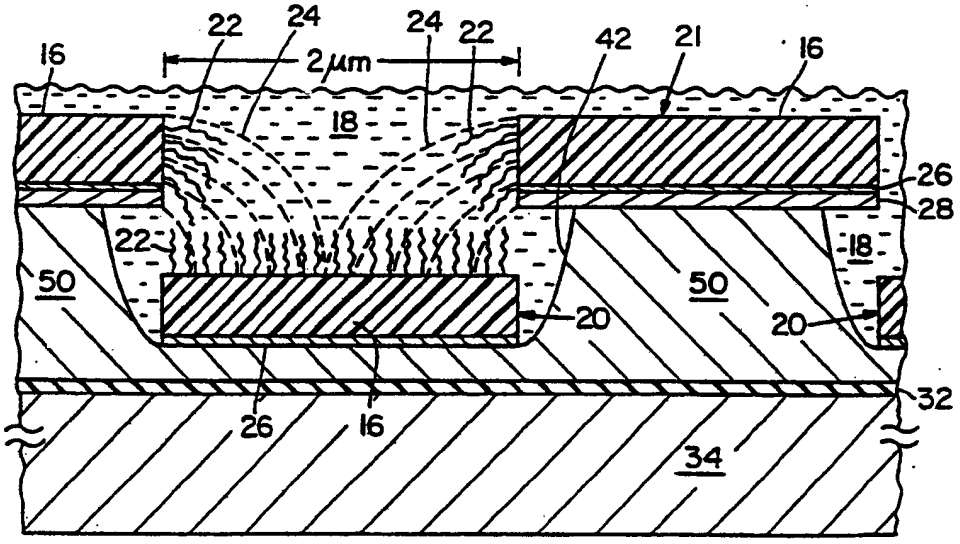

(57) Abstract

A method and apparatus are disclosed for identifying molecular structures within a sample substance using a monolithic array of test sites formed on a substrate upon which the sample substance is applied. Each test site includes probes formed therein to bond with a predetermined target molecular structure or structures. A signal is applied to the test sites and certain electrical, mechanical and/or optical properties of the test sites are detected to determine which probes have bonded to an associated target molecular structure.

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must be implemented when using radioactive markers or labels to reduce the exposure to radioactivity. Typically, workers must wear a device to continually monitor radioactive exposure. In addition, pregnant females should take additional precautions to prevent the occurrence of genetic mutations in the unborn.

The conventional radioactive detection scheme has sensitivity limitations in both the temporal and spatial domains. The use of radioactive labelling currently has a spatial resolution of one millimeter. Additional hardware and software are required to reduce the spatial resolution below one millimeter.

The sensitivity of detection utilizing autoradiographic film is directly related to the amount of time during which the radioactive labelled fragments are exposed to the film. Thus, the exposure time of the film may range from hours to days, depending upon the level of radioactivity within each detection test site. A β scanner may drastically reduce the time required for film exposure during radiography. However, the use of the β scanner significantly increases the expense associated with this type of detection, and has intrinsically poor spatial resolution.

Optical detection of fluorescent labelled receptors has also been utilized to detect molecular binding. Briefly, for DNA sequence analysis applications, a base-specific fluorescent dye is attached covalently to the oligonucleotide primers or to the chain terminating dideoxynucleotides used in conjunction with DNA polymerase. The appropriate absorption wavelength for each dye is chosen and used to excite the dye. If the

OPTICAL AND ELECTRICAL METHODS AND APPARATUS FOR MOLECULE DETECTION

Description

Background of the Invention

In many applications, it is desirable to detect the presence of one or more molecular structures in a sample. The molecular structures typically comprise ligands, such as, cells, antibodies and anti-antibodies. Ligands are molecules which are recognized by a particular receptor. Ligands may include, without limitation, agonists and antagonists for cell membrane receptors, toxins, venoms, oligo-saccharides, proteins, bacteria, and monoclonal antibodies. For example, a DNA or RNA sequence analysis is very useful in genetic and disease diagnosis, toxicology testing, genetic research, agriculture and pharmaceutical development. Likewise, cell and antibody detection is important in disease diagnosis.

A number of techniques have been developed for molecular structure detection. In DNA and RNA sequence detection, two procedures are generally used, autoradiography and optical detection. Autoradiography is performed using ^{32}P or ^{35}S. For DNA sequence analysis applications, nucleic acid fragments are end labeled with ^{32}P. These end labeled fragments are separated by size, then exposed to x-ray film for a specified amount of time. The amount of film exposure is directly related to the amount of radioactivity adjacent to a region of film.

The use of any radioactive label is associated with several disadvantages. First, prolonged exposure to radioactive elements increases the risk of acquiring genetic diseases, such as cancer. As such, precautions

absorption spectra of the dyes are close to each other, a specific wavelength can be chosen to excite the entire set of dyes.

A particular optical detection technique involves the use of a dye, for example, ethidium bromide, which stains duplexed nucleic acids. The fluorescence of these dyes exhibits an approximate 20-fold increase when it is bound to duplexed DNA or RNA, when compared to the fluorescence exhibited by unbound dye, or dye bound to single-stranded DNA. This type of dye is used to detect the presence of hybridized DNA (or RNA) during a hybridization experiment. Although the use of conventional optical detection methods increases the throughput of the sequencing experiments, the disadvantages are serious.

Therefore, a need has arisen in the industry for a safe, low-cost, fast and accurate method and apparatus for detecting molecular structures at reduced complexity.

SUMMARY OF THE INVENTION

In accordance with the present invention, a method and apparatus for detecting the presence of molecular structures in predetermined test sites is provided which substantially eliminates or prevents the disadvantages and problems associated with prior devices.

In an electrical embodiment of the present invention, a substance having a molecular structure is applied to a plurality of test sites, each test site having a probe formed therein capable of binding to a known molecular structure. Electrical signals are

applied to the test sites, and electrical properties of the test sites are detected to determine whether the probe has bonded (hybridized) to, or with, an associated molecular structure.

The test sites are monolithic structures formed on, or in, semiconductor chips or wafers using very large scale integrated (VSLI) circuit methods. This results in a low-cost, small-size, testing device which may be inexpensive enough to be disposable after use.

Hybridized molecules can be detected, in accordance with one embodiment of the invention, by sensing the change in dissipation of a capacitor formed at the test site, or by sensing the change in AC conductance of a test site when hybridized molecules are present. Alternatively, by forming a transmission line between two electrodes at each test site, the presence of hybridized molecules can be detected by measuring the RF loss associated with the formation of hybridized molecules at the test site.

In another embodiment, micro-machined resonators are formed in each test site and the change in resonant frequency, or the change in the Quality Factor (Q) of the resonator, caused by formation of hybridized molecules may be measured to determine which sites contain hybridized molecules.

In an alternate optical embodiment of the invention, a charge-coupled-device (CCD) array is provided, with each electrode of the CCD array aligned with a respective adjacent test site. Light attenuation, caused by greater absorption of illuminating light in test sites with hybridized molecules is used to determine the sites with the

hybridized molecules. The CCD array can be integrated with a corresponding test site array. Alternatively the test site array may be a separate disposable plate.

The probes within each test site are all identical, but differ from test site to test site. The probes for DNA or RNA sequence testing are generally formed of oligonucleotide strands. In accordance with another embodiment of the invention, an optical direct patterning system is used to perform localized sensitization of the microarray or localized synthesis of oligonucleotide strands at each test site to customize or differentiate each of the probe strands.

A further understanding of the nature and advantages of the invention herein may be realized with respect to the detailed description which follows and the drawings described below.

Brief Description of the Drawings

Fig. 1 is a schematic partial perspective of a microelectronic sensor array in accordance with a preferred embodiment of the invention.

Fig. 2 is an enlarged view of a portion of Fig. 1.

Fig. 3 is an enlarged view of the electrode portion of Fig. 2.

Fig. 4 is a section taken along lines IV-IV of Fig. 3.

Figs. 5A-5D are schematic cross-sectional process diagrams showing important steps in forming test sites.

Figs. 6A-6H are schematic cross-sectional process diagrams showing important steps in forming alternate embodiments of the test sites.

invention may also be used for cell detection and antibody detection or detection of any hybridized molecule.

The sequencer 10 comprises an X-Y array of test sites 12 electronically addressable by conductive leads X1, X2, X3---XN on the X-axis and conductive leads Y1, Y2, Y3---YN on the Y-axis. X-logic circuitry 36 for sequentially addressing each X-line is coupled to detection and recognition circuitry 40. Similar circuits 56 are coupled to the Y-lines Y1---YN. The array 10 and X and Y logic circuitry 36 and 56 and circuitry 40 may all be implemented on a single semiconductor chip depending upon cost trade-offs.

The test sites 12, described in greater detail hereinbelow, are formed in a semiconductor wafer using semiconductor photolithographic processing techniques. Each test site contains a plurality of probes 22 (See Fig. 4) which are capable of binding to known molecular structures (hereinafter "target(s)"). The targets could comprise, for example, biopolymers such as polynucleotides, DNA, RNA, cells, antibodies or anti-antibodies. For the case of a RNA or DNA sequencer, the synthetic probes may comprise, for example, oligonucleotides. All the probes 22 in a given test site are identical. But, the probes in respective test sites 12 differ in a known sequence for simultaneous detection of a plurality of different targets (or subsequences within a target molecule) within a single array 10.

When a sample substance containing the targets in an electrolyte solution 18 is poured onto the array 10, the targets bind with associated probes 22 within a

plurality of wells 42 formed in each test site 12. After sufficient time for binding, the surface of the array 10 is rinsed to remove excess targets or other unbound molecular structures. The remaining target structures will be, for the most part, bound to the probes attached to the microfabricated array 10 at specific test sites 12. Each test site 12 is then interrogated electronically by the logic circuitry 36 and 56 to determine whether targets have bound in that test site. Test sites having bound targets, i.e., hybridized molecules, will have changed electrical parameters, which may be detected by detection circuitry 40 coupled to the test sites over the X and Y leads. Thus, by electronic addressing, the detection of specific target/probe bindings is achieved at each test site 12 within the microfabricated array 10, thereby determining the composition of the targets that remain present after washing.

For the example of DNA sequencing, recognition circuit 40 performs a sequence analysis described in connection with Fig. 21 based upon the composition of the targets (nucleic acids) detected by the circuitry 40.

Note: Circuit 40 is preferably coupled to the test sites by transistor switches (not shown) using row and column addressing techniques employed, for example, in addressing dynamic random access memory (DRAM) or active matrix liquid crystal display (AMLCD) devices.

rinse, a number of the wells in the array will contain a significant amount of this bonded or hybridized DNA, and the rest will contain only the original oligonucleotide strands in an aqueous solution. The wells are then interrogated electrically in sequence using the electrodes 16 and 20 in each site. The sites that contain hybridized DNA are recorded. For example, sites without hybridized DNA will have different electrical properties than those with hybridized DNA and will not be recorded. At the resonant frequency of a DNA molecule in aqueous solution, the imaginary part ϵ'' of the complex relative permittivity $\epsilon_r = \epsilon' - j\epsilon''$ of the solution can be approximately a factor of 10 to 100 times larger than its value for an aqueous solution without the DNA. Methods B, C, D, and E below are designed to measure or detect this difference in ϵ'' at each site 12. From this data base, a computer "overlapping" or "neural network" algorithm in circuit 40 reconstructs the entire coding sequence of the unknown DNA.

B. Dissipation Factor Test

Fig. 7 is a plot of dissipation factor versus the log of frequency for bonded (hybridized) DNA (curve B) and unbonded DNA (curve A) showing how the dispersion factor $D = \epsilon''/\epsilon'$ differs, depending upon whether the DNA is bonded or not. Note: Depending upon the particular samples measured, the curves of Fig. 7 may be reversed, i.e. curve B could represent unbonded DNA. This difference in dispersion factor is used to determine the

presence or absence of hybridized DNA at a test site formed as in Figs. 1-6. The dissipation factor at each test site is measured by well-known instrumentation such as an LCR meter in circuit 40. The meter is successively coupled to each site 12 via logic circuits 36 and 56.

C. AC Conductance Test

Similarly, the presence or absence of hybridized DNA can be detected by measuring the AC conductance G_{AC} = ϵ"A/d at each test site; wherein A is the effective area of one electrode and d is the effective distance between electrodes. At the relaxation frequency of a given DNA molecule, the AC conductance should be as much as 100 times or more larger than the conductance when no DNA is present. Fig. 9 is a schematic representation of how this test may be conducted. A pulsed or frequency-scanned waveform is applied across electrodes 21B and 20B of each test site 12B. Probes 22 are formed on each electrode and an aqueous solution of target molecules is formed in the wells 42B of the test sites 12B. The presence of hybridized DNA is detected at a resonant frequency of DNA as shown in Fig. 10.

An LCR meter may be used to measure G or R = 1/G at a discrete frequency. Alternatively, as discussed in connection with Figs. 9 and 10, G can be measured as a function of frequency.

D. Transmission-Loss Detection Test

Signal loss on a transmission line is also sensitive to ϵ''. By incorporating a transmission line 11 between the X and Y lines at each test site (as shown in Fig. 8) electrical detection of hybridized molecules, such as DNA, can be accomplished by scalar measurement of the RF loss of an electromagnetic wave passing along the line 11 at each test site 12A. Line 11 may comprise a micro-miniature version of stripline, microstrip, waveguide, coplanar waveguide, slotline, or coaxial line. To achieve maximum sensitivity with this method, the test site well 42A is made relatively wider and/or longer than the wells in Fig. 4, and the length of the transmission line in the well is maximized by forming it in a meandering pattern.

E. Pulse and Chirp Method of Detection

As shown in Fig. 11, a frequency scanned or chirped voltage waveform V_i may be applied across the electrodes at each site and the resultant response waveform V_o (Fig. 12 or Fig. 13, depending upon whether frequency is increasing or decreasing) is analyzed to determine the presence of hybridized DNA as indicated by a maxima at a hybridized DNA frequency. The measurement of the relaxation frequency of the hybridized DNA using a frequency-scanned waveform gives additional information about the properties of the hybridized DNA, e.g., crosslinked versus non-crosslinked.

F. Micromechanical Resonator Detection Methods

In this embodiment, a plurality of mechanical resonator structures are formed in test sites formed in silicon wafer 34C, as shown in Fig. 14. The resonator structure comprises a lower metal sensor electrode 20C extending in the X-direction and an upper membrane resonator film 21 preferably of silicon nitride or metal such as tantalum extending along a Y-direction in the plane of the wafer. Typically the membrane size is about 100 microns in diameter or width/length. A dielectric gap 60, preferably of air, is formed between the upper and lower members 21C and 20C.

A test site well 42C is formed over membrane 16C and probes 22C formed in the well surfaces. Target DNA solution 18C is dispensed into the test well 42C. The mechanical cavity 60 between the upper and lower electrodes 16C and 20C forms a resonator. This resonator has a resonant frequency in the kilohertz to multimegahertz range with a narrow resonant linewidth.

An RF signal propagated across each resonator will produce a characteristic high Q response with a narrow linewidth. A shift in either Q or resonant frequency indicates the presence of hybridized molecules on the resonator surface electrode membrane 21C.

Membrane electrode 21C may be formed of a thin film of silicon nitride using chemical vapor deposition at a well controlled silicon to nitrogen ratio and controlled elevated temperature to adjust the film tension when it is cooled to room temperature. Membranes can be formed on unpatterned silicon wafers then released as free standing structures by etching out a silicon window from the back side. Examples of mechanical resonators and

The advantages for using 1,2-dioxetanes are numerous. In addition to no radioactivity exposure, this method is relatively simple to perform (reagents and equipment are inexpensive). Finally, this method has a low background noise level and wide dynamic range.

In an alternative two-piece implementation as shown in Fig. 16 the probe site array 200' is formed on a separate thin transparent substrate such as a 10-mil-thick pyrex plate 270. This separate plate is marked with precision alignment features such as etched or printed gratings (not shown) to permit a precise automated overlay of the separated probe plate onto a separated CCD array 260. Each array location in the probe plate is sensitized with unique probes. The CCD array is then fabricated with or without the blocking filter 250 of Fig. 15. In one embodiment, an analysis is made by bringing the probe plate into registered close proximity over the CCD array without using a lens to image the plate onto the CCD. Irradiation of the plate is as in either of the embodiments discussed above in connection with Fig. 15. A further alternative is to image the separated probe plate 200' onto the CCD array 260 using a lens. This would allow a greater separation between the plate and the CCD array, for the case in which secondary fluorescence is used, and also allows separation of the excitation and fluorescence by obliquely exciting the probe plate. Imaging with magnification or demagnification is possible so that the probe plate dimensions can be optimized separately from the CCD.

The CCD device used to monitor the probe array for any of these geometries can be of the conventional variety and sensitive to the ultraviolet and visible spectrum. An alternative approach is to use an infrared, heat-sensitive array detector such as a platinum silicide or iridium silicide infrared imager. This latter choice would permit the direct monitoring of heat evolved from the probe array during a biochemical reaction such as hybridization or antibody action. DNA hybridization and other heat-generating reactions may be directly detectable through their thermal signature during reaction. The infrared transmission and reflection properties of the product (e.g., hybridized DNA) will be distinctly different than the reactants due to the formation of new molecular bonds with new absorptions from infrared-active vibrational and rotational modes in the product molecule. In the configuration of Figs. 15 and 16, thermal properties can be monitored also by monitoring thermally generated noise in a conventional visible wavelength or IR detector array. In this case heat generated by the biochemical reaction is transmitted by thermal conduction through the thin device layers and detected as a noise burst on the electrode 220. The array may also be flood-irradiated with infrared, visible, or ultraviolet light in the configuration of Fig. 15. In this case, light is chosen specifically in a product-state (e.g., hybridized DNA) absorption band. In the unreacted state the flood illumination is transmitted through the well and reflected by filter 250. Wells in which the desired reaction has occurred become absorbing

at the flood illumination wavelength. After absorption the flood illumination automatically converts to heat and is detected after conduction into the device below the active well site.

V. PROBE FORMATION

A. General

One method of forming the array 10 uses probes attached to the test sites 12 in the array. Different probes can be attached to the test sites 12 according to the type of target desired. Oligonucleotides, single or double stranded DNA or RNA, antibodies or antigen-antibody complexes, tumor cells and other test probes known to those of skill in the art may be used. The probes are attached to the test sites by fixation to a solid support substrate on the surface of the wells 42, or alternatively, attached directly to the electrodes 16 or 20, as in Fig. 4. The solid support substrates which can be used to form the surface of the wells 42 include organic or inorganic substrates, such as glass, polystyrenes, polyimides, silicon dioxide, and silicon nitride.

The solid support substrates or the electrodes must be functionalized to create a surface chemistry conducive to the formation of covalent linkages with the selected probes. As an example, a glass support can be functionalized with an epoxide group by reaction with an epoxy silane. The epoxide group on the support reacts with a 5'-amino-derivatized oligonucleotide probe to form a secondary amine linkage, as described in Parkam and Loudon, BBRC 1:1-6 (1978), which is incorporated by

reference herein. Formation of this covalent linkage attaches the probes 26 to the support surface in the desired array. Examples of functionalized polystyrene surfaces include 5′ aldehyde or carboxylic acid derivatives coupled to hydrazide-activated polystyrene, as described in Kremsky, et al. (1987) Nucl. Acids Res. 15:2891-2909, and 5′ amino derivatives coupled to polystyrene which has been activated by diazotization and 5′ phosphate derivatives coupled to amino-functionalized polystyrene, as described in Lund, et al. (1988) Nucl. Acids Res. 16:10861-10880, both articles being incorporated by reference herein.

For direct attachment of probes to the electrodes, the electrode surface must be fabricated with materials capable of forming conjugates with the probes. Materials which can be incorporated into the surface of the electrodes to provide for direct attachment of probes include electrometal materials, such as gold, niobium oxide, iridium oxide, platinum, titanium, tantalum, tungsten and other metals. These electrometals are capable of forming stable conjugates directly on the plate surface by linkages with organic thiol groups incorporated into the probe, as described in Whitesides et al. (1990) Langmiur 6:87-96 and Hickman et al. (1991) J. Am. Chem. Soc. 113:1128-1132, both of which are incorporated by reference herein. As an example, a synthetic DNA probe labeled with a thiol group at either the 5′ or 3′ end will form a stable conjugate with a metal, such as gold, in the plate surface to create an array of directly attached probes.

Laser initiation of the reaction occurs either by localized heating or by photochemistry. A preferred embodiment uses a visible-wavelength or UV argon ion laser in combination with a galvanometer scanning system to initiate photochemical synthesis. Alternatively, since synthesis reactions are known to be highly temperature sensitive, an argon or infrared laser may be used to initiate synthesis by local heating of an array site.

The method can also be applied to the synthesis of peptides or other polymeric probes on solid supports, based upon the principle of thermally addressable de-protection. For example, in the case of peptide synthesis, site selective peptide synthesis is achieved by thermal removal of the f-moc protecting groups, typically in dilute base, followed by capping and the other ordinary steps of peptide synthesis.

Alternatively, a "glue" layer can be locally activated Fig. 26A-D (or deactivated) or locally applied Fig. 25A-D to a test site by means of scanned laser irradiation. In this embodiment the ultraviolet, visible or infrared laser is used to photochemically or thermally alter the adhesion properties of the desired array sites. The probe solution, for example of type A, is then washed over the array resulting in localized adhesion of the type A probe at the desired sites. The type A probe solution is then rapidly rinsed from the system, a second laser irradiation at new array sites is applied, and type B probe solution is introduced to adhere type B probes. The process is repeated to sensitize the full array.

Array sensitization may be accomplished using, for example, a CW argon-ion or CW Nd:YAG laser using scanning optics such as galvanometers or rotating mirrors, or using a fixed laser beam with a computer-controlled X-Y stage. An activation or deactivation process in a "glue" layer can be preferably accomplished using a short-pulsed laser such as a pulsed Nd:YAG laser or excimer laser. An excellent approach is to simply cover the "glue" layer 902 to "deprotect" and thereby reveal the "glue" by ablating a passivating material 904 applied over the "glue" (See Figs. 26A-D). Examples of "glue" layers are epoxides, thiols or hydrophilic, e.g., hydrated surfaces. Passivating materials can be hydrophobic materials such as fluorine-terminated fluorocarbons or the derivatives or hexamethyldisilizane.

Figs. 25A-D and 26A-D illustrate two alternate methods of probe formation using the "glue" approach. Furthermore each show two alternate ways to activate a test site. One way is to use a programmable element such as a heater element 906 embedded beneath a test site to induce a thermal reaction in the test site and thereby create or deposit a glue layer 920 to which the probes adhere. Fully synthesized probes 912 are washed over the cite and adhere to the exposed glue layer site 920, Fig. 25D. Next another site is formed or exposed and a different probe attached. Alternatively external radiation as in Fig. 25B is used to form the glue layer 920; or as in Fig. 26B and C to ablate a passivating layer 904 and expose a glue layer 902.

the washing (removal) of unhybridized and mismatched target molecules. This technique is not only applicable to the electronic genosensors of Figs. 1 through 9, which have electrodes present within each test site, but can be employed in both the micromechanical-resonator and CCD-based approaches by either using the electrodes present within or under each test site or fabricating one or more additional electrodes at each test site for this purpose.

Alternatively, the potential applied to individual wells can be used to draw a current surge through the well structure sufficient to evaporate a "glue" layer or glue passivating layer similar to that described above in the last method. Sensitization of the array is similar to the electronic programming of an array of electrical fuses.

Referring now to Figs. 18 and 19, a microfluidic system for synthesizing unique genosensor probes in situ in a test site will now be described. In this embodiment, reagent sources 352 are individually fluidly coupled via channels L1, L2 --- LN to respective microchannel valves V1,V2 --VN formed in a suitable substrate 341. Valves V1-VN enable flow of solution into manifold line L4. Microfluidic peristaltic pump P1 forces the solution onto array 10', which is enclosed by laser-radiation-permeable films 344 and 343, such as silicon nitride or silicon dioxide.

Radiation from laser 416' is selectively projected onto individual test sites 12' formed in substrate 341, in accordance with previously described scanning or imaging methods. Laser scanning of test sites induces

localized activation of individual sites as the input solution fluids are rapidly switched using valves V1-VN.

The entire fluidic system as well as the array may be formed on a single chip of semiconductor or dielectric material, such as Si, glass, Al_2O_3, etc. Channels 342 are etched into the substrate 341 using conventional photolithography and etchants or by micro-machining techniques. An array 10' of test sites 12' is formed in the substrate, as described in connection with Figs. 1-6.

The microfluidic flow system depicted in Figure 19 can be formed as follows. A photoresist material is spin-coated on a substrate 341, formed, for example of pyrex glass. The microchannel structure is then patterned into the photoresist using standard photolithography and the pattern, including channel structures 343 and 342, are transferred into the substrate by etching using buffered HF. A membrane actuator layer 344, comprised of preferably a piezoelectric, such as lead zirconium titanate or PVDF polymer and metal electrodes, is then bonded to the microchannel structure. During sensitization the array 10' is sealed against the microfluidic system preferably using an elastomer O-ring 345. Alternate membrane actuator layers, known to specialists in the art, make use of shaped memory alloys rather than piezoelectrics, or are based on passive materials deformed electrostatically, for example, aluminum films which are deflected by DC voltages applied to electrodes (not shown).

The concept of using a larger set of DNA probes to decipher the base sequence of a DNA sample target is illustrated below. Example I shows a small portion of the base sequence of a DNA sample, which has been converted to single-stranded form by heating prior to analysis. By exposing the sample DNA to a set of synthetic DNA probes representing all possible sequences for a given probe length (for example, all 65,536 8-base probes), and then detecting which probes have specifically bound to the target DNA, a complete list of oligonucleotide sequences contained in the DNA sample can be generated. In the case shown in Example 2 (below) only those 8mer probes listed would hybridize to the sample DNA sequence. In turn, an overlapping algorithm is used to generate the complete sequence of the target DNA from the oligonucleotide content.

Example I

Unknown Single Strand DNA (Target)

ATCGCTTACGGTAATC

Example II

Hybridized Synthetic Genetic Probes

TAGCGAAT
AGCGAATG
GCGAATGC
CGAATGCC
GAATGCCA
AATGCCAT
ATGCCATT
TGCCATTA
GCCATTAC

VIII. APPLICATIONS

Commercial applications of the present invention with regard to DNA and RNA detection include genetic research, genetic and infectious disease diagnosis, toxicology testing, individual identification, agriculture identification and breed optimization, quality assurance through contaminant detection, and occupational hazard screening via mutation detection.

There are currently estimated to be 4,000 to 5,000 genetic diseases in humans, in which a mutational change in a gene destroys or hinders the function of a gene product, leading to a serious medical condition. The affected genes and proteins (gene products) have thus far been identified for a small fraction of human genetic diseases, although the number is increasing steadily. A few examples of human genetic diseases for which mutations associated with the disease have been identified include cystic fibrosis, phenylketonuria, Alzheimers' disease, cancer, Duchenne muscular dystrophy, and familial hypercholesterolemia. Although, in some cases, the disease is associated with one or very few specific mutations, it is becoming evident that many, if not most, genetic diseases can be caused by any of numerous mutations, scattered along the affected gene. In the former case, the presence of a defective gene can be detected through the use of simple DNA hybridization detection tests in which a synthetic DNA probe is used to discriminate between a wild type and mutant DNA sequence. In the latter case, a substantial DNA sequencing effort is required to search through an entire gene for mutations that may be associated with a disease.

(RFLP) analysis, which is time consuming and laborious. DNA typing can play an important role in forensics and paternity testing. In addition, there is interest in DNA typing all personnel in the armed services.

As valuable new plants and livestock are developed by genetic engineering, there will be a need for DNA typing to verify the source and ownership of agricultural products. The sequence information that will come from genome sequencing in humans, plants and animals will lead to increased application of genetic engineering techniques to develop pharmaceutical agents and create improved crops and livestock. Examples include strains that are more resistant to disease and harsh climates, as well as crops that have a greater yield or higher nutritive value.

The present invention can be used in connection with detection of targets which are molecular structures other than DNA or RNA, such as cells and antibodies. Table III sets forth feasible probe types for other molecular structures serving as targets. The stated probe types are not meant to be exclusive.

TABLE III

Probe Types

Target	Probe
DNA, RNA	Oligonucleotide
Antibody	Antigen (peptide), anti-antibody
Cell	Antibody, protein
Hormone receptor	Hormone
Aviden	Biotin
Immunoglobulin	Protein A
Enzyme	Enzyme Factor
Lectins	Specific Carbohydrate

When the detector employs peptides or other antigens as probes, it can be used to detect antibodies in biological fluids, as shown in Fig. 22.

In this embodiment, a peptide antigen (the probe 22) is affixed to the SiO_2 50 at the bottom of the test well 12A (similar to that illustrated in Fig. 6H), employing a bifunctional crosslinker such as one with a silane at one end and an epoxide or other peptide specific group at the other.

The treated surface is then incubated with a fluid 18 containing antibody (the target T). Because antibodies are large macromolecules (150,000 to 950,000 MW, depending on class), the resulting target/probe bonding produces a large change in the permittivity of the test well 12A. The magnitude of the effect can be additionally amplified by treating the target/probe complex with a second antibody which is specific for the target antibody, thereby creating a very large complex.

The affinity and selectivity of antibody/antigen and antibody-antibody interaction are well known and are the basis for an existing class of biotechnology (ELISA assays, immunohistochemistry, and others). The technology described here employs those well understood binding interactions in a new microelectronic detection scheme.

The commercial application of the methodology is for use to detect the presence of any of hundreds of thousands of different antibodies or other proteins, simultaneously, in a blood sample or other biological fluid. This is particularly useful in blood typing, the

detection of viral infection such as AIDS, or the diagnosis of cancer. It would also be very useful as a research tool. It would replace or augment the use of ELISA assays and other biochemical methods to detect antibody/antigen interaction.

When the detector employs as a probe, peptides, antibodies or other molecules which bind to cells, it can be used to detect specific cell types in biological fluids.

In this embodiment, the probe 22 comprises an antibody, protein or other molecule which is known to bind to the cell surface. The target T in this case is an intact cell having receptors T for bonding with the probes 22.

A fluid solution containing cells is added to the detector. Subsequent to the target/probe binding interaction, binding gives rise to detector wells which are coupled to a cell. Since cells do not conduct current and display low frequency dielectric relaxation, binding of a cell can be detected by either a change in absolute conduction in a well (a modification of the Coulter principle) or by the induction of a low frequency dielectric relaxation effect.

The commercial application of the methodology is for use to detect the presence of cells with altered cell surface properties, especially cells in the blood or other bodily fluids. Cells from solid tissues could be analyzed subsequent to standard tissue dispersement methods. Such a detector would be useful in the diagnosis of viral infection and for cancer diagnosis, as well as a scientific research tool. It would serve

as a replacement for the use of fluorescence microscopy (immunohistochemistry) and fluorescence activated cell sorting.

IX. ADVANTAGES

Current microfabrication techniques enable inexpensive construction of multimegabit memories that exhibit uniform densities and properties. Hence arrays containing potentially millions of individual biological test wells or sites can be miniaturized comparable to standard electronic devices at a similar cost. For example, a 1cm by 1cm array could easily be fabricated containing one million biological test sites. Moreover, the uniform electrical properties of the devices fabricated in such manner enhance the detection sensitivity beyond many other approaches.

One important advantage of the microfabricated electronic detector and the optical-absorption CCD detector described previously is that the detection method provides direct detection of target/probe molecular binding. Hence no toxic fluorescent, radioactive, or chemical marker need be attached to the targets or probes. Rather, only an appropriate electrical signal or frequency shift must be experienced for detection. Such signals or shifts naturally occur for many target/probe combinations, such as DNA and RNA to an oligonucleotide. However, if the signal or shift in the electronic detector is weak or nonexistent after bonding, a charged molecular marker can be attached to the target. In addition, detection in the electronic detector is observed by a change in frequency

characteristics, as opposed to a change in magnitude characteristics which can be obscured in time as the microfabricated array is exposed to the corrosive biological solutions. Thus, the device may be cleaned and reused a number of times without affecting its accuracy. Although the method of detection will withstand some corrosion of the electrodes, a passivation layer can be employed to coat the plates for even longer use.

Another advantage of the present invention is that the electronic circuitry used to interrogate the test sites to perform the detection measurements can be fabricated directly on the wafer containing the biological array. Switch matrices, signal processing circuitry, and energy sources could all be incorporated on the same chip to facilitate rapid detection across the array. Consequently, the incorporation of active circuitry on the wafer would also greatly reduce the cost of experimentation.

The density of the probes 22 attached at the test site 12 directly determines the sensitivity. The microelectronic method has been shown to provide a factor of ten discrimination between short (nonhybridized) and long (hybridized) single-stranded DNA fragments, whereas the intercalating-dye optical approach provides a factor of three.

The elimination in most embodiments of radiographic film reduces the testing time since film exposure is not required. Sample preparation time is reduced greatly since the nucleic acid fragments need not be labeled. The detection method is quick; the measurements can be

performed as soon as sufficient molecular binding is completed. Furthermore, the measurement process can be automated via on-chip microprocessor control to provide a very fast method of accessing each test site in the array.

The microelectronic technology incorporated into these types of detection devices will drastically reduce the price for such experimentation. Essentially, the efficient mass production techniques employed in making megabit memory chips and megapixel CCD imaging chips can be employed.

Although the present invention and its advantages have been described in detail, it should be understood that various changes, substitutions and alterations can be made herein without departing from the spirit and scope of the invention as defined by the appended claims.

For example, the active circuitry of the genosensor array, such as circuits 36, 56, 38, 58 and 40 of Fig. 1, can be integrated monolithically with the array of wells or the same substrate. Switch matrices, analog testing circuits, and analog or digital (microprocessor) controllers could all be fabricated on the same wafer to perform or simplify the electrical tests. As shown in Fig. 24, transistors, such as, TRX 1, could be integrated into each substrate adjacent to a respective test site 12, for example, to disconnect each site electrically, except when it is being sampled. This would necessitate an additional address line A3 for each column but would reduce parasitic capacitance and spurious signals from lines not in use. A greater

reduction of these undesired effects could be achieved by a second address line and set of transistors coupled to the Y-side of the site 12.

CCD circuitry (including CCD implementations of neural networks) has been demonstrated that can perform a wide variety of signal processing and pattern recognition functions. Integration of a CCD data-processing circuit with a genosensor array could simplify the DNA detection and decoding, and would be compatible with the integrated CCD imager, as described in connection with Figs. 15 and 16.

While the invention has been illustrated in connection with a wet type of testing in which solutions are used; it is entirely feasible to use a "dry" or "gel" approach in which the probes and hybridized probe/target combinations are formed in a dry medium or in a gel.

CLAIMS

1. A method for identifying molecular structures comprising the steps of:
   a) forming an array of test sites, each site having probes formed therein capable of bonding with a unique molecular structure and wherein the probes in each test site differ from the probes in other test sites;
   b) applying a sample substance to the test sites;
   c) applying a test signal to the test sites; and
   d) detecting properties of the test sites resulting from the applied signal to determine which probes have bonded to a molecular structure in the sample substance such that a plurality of molecular structures can be identified.

2. The method of Claim 1 wherein the test signal is an electro-magnetic signal and forming said array comprises the steps of:
   a) forming a first layer on a substrate;
   b) forming a second layer on the first layer;
   c) forming openings in the second layer into the first layer exposing a portion of the first layer; and
   d) forming a pair of electrodes in the openings and wherein said test signal is applied to the electrodes.

3. The method of Claim 2 wherein forming said pair of electrodes comprises the steps of depositing metallization on the second layer after the openings are formed; which metallization forms an upper electrode on the surface of the second layer between openings and a lower electrode on the exposed portions of the first layer.

4. The method of Claim 3 wherein the substrate is formed of silicon, and the first layer and second layer are formed of a silicon based dielectric.

5. The method of Claim 4 wherein the first and second layers are SiO_2 and Si_3N_4, respectively, and the metallization comprises Al, Ti, Pt, W, Ta and their silicides or Au.

6. The method of Claim 1 wherein said step of detecting comprises detecting the dielectric properties of the test site.

7. The method of Claim 1 wherein said step of applying an electronic signal comprises applying a pulsed or varying frequency signal.

8. The method of Claim 1 wherein each test site is formed with a resonant structure which is resonant in a frequency range of electrical signals.

9. The method of Claim 8 wherein said detecting step comprises detecting changes in the Q or the resonant frequency of the resonant structure.

10. The method of Claim 1 wherein said sample substance is in a solution or a gel.

11. Apparatus for identifying molecular structures within a sample substance, comprising:

a) test sites formed on a substrate for receiving the sample substance, each test site having row and column electrodes formed therein and the structure having row and column leads extending to a respective electrode at each site;

b) probes formed in said test sites for binding to molecular structures;

c) circuitry for applying an electronic signal to the electrodes of the test sites; and

d) circuitry for detecting electrical properties of the test sites to determine which probes have bonded to a molecular structure in the sample substance.

12. The apparatus of Claim 11 including an array of resistors formed beneath the test sites.

13. The apparatus of Claim 11 wherein said electrodes comprise a plurality of conductive fingers extending from a base.

14. The apparatus of Claim 13 wherein the spacing between said fingers is less than about 30 microns.

15. The apparatus of Claim 14 wherein a first of said plurality of electrode fingers is disposed in a lower portion of a plurality of wells formed in said substrate and a second of said plurality of electrode fingers is disposed on said substrate above the fingers on the lower portion.

16. The apparatus of Claim 11 wherein said probes comprise molecular probes from the group comprising cell probes, antibody probes, or peptide probes.

17. Apparatus for determining the presence of molecular structures within a sample substance comprising:

a) a test site array formed on a substrate for receiving the sample substance;

b) probes formed in the test sites for binding to molecular structures; and

c) a detector array having a plurality of detectors, each respective detector disposed adjacent to a respective test site and wherein radiation is propagated through said test sites and is absorbed to a different degree by sites containing bonded probes than non-bonded probe test sites and such difference is sensed by the detectors and used to generate a signal to identify the presence of the molecular structures within the sample substance.

18. The apparatus of Claim 17 wherein said test site array is formed of a disposable plate which is separable from the detector array.

19. The apparatus of Claim 17 wherein the test site array is formed integral with the detector array to form an integrated structure.

20. The apparatus of Claim 18 wherein the disposable plate is formed of quartz, glass, plastic, Al_2O_3 or polyimide.

21. The apparatus of Claim 11 wherein the electrodes in each test site are coupled together by a transmission line.

22. A circuit for determining the presence of molecular structures within a sample substance, comprising:

a) a substrate;

b) a plurality of test sites formed in said substrate;

c) electrodes formed in each of the test sites;

d) leads extending to each of the electrodes;

e) probes formed in respective test sites, said probes of each respective test site being identical in structure, and probes of different test sites being of various structures in order to bond with respective predetermined molecular structures.

23. The apparatus of Claim 11 further including an address lead coupled to one of said electrodes via a transistor switch.

24. The apparatus of Claim 17 wherein the radiation is generated by radioactive, luminescent, or chemiluminescent labels within the test sites.

25. The apparatus of Claim 17 wherein the radiation is generated by secondary emission stimulated by photon irradiation of the test sites.

26. The apparatus of Claim 17 wherein the radiation is infrared radiation and the detectors sense thermal energy.

27. Apparatus for synthesizing molecular probes in situ comprising:

a) an array of test sites each site containing molecules to be synthesized;

b) a light source for irradiating selected sights with light to induce synthesis of molecules in said selected site.

28. The apparatus of Claim 27 wherein the light is in the visible wavelength range and photochemical synthesis occurs.

29. The apparatus of Claim 27 wherein the light source is a laser which is scanned from site to site to induce localized synthesis.

30. The apparatus of Claim 27 wherein the light induces local heating of a selected test site to induce a thermal synthesis of molecules.

31. The apparatus of Claim 27 wherein the molecules comprise oligonucleotide strands.

32. Apparatus for synthesizing molecular probes in situ comprising:

a) an array of test sites each site containing precursor molecules to be reacted;

b) an array of resistors disposed adjacent to the test site array with a respective resistor located in proximity to a respective test site; and

c) coupling means for coupling a respective resistor to an electrical source for heating a respective resistor to induce a thermal reaction for synthesis of molecules in a respective test site.

33. The apparatus of Claim 1 wherein the test site array and resistor array are formed as an integrated structure.

34. Apparatus for synthesizing molecular structures in situ comprising:

a) an array of test sites each site containing precursor molecules to be reacted; and

b) each test site containing electrodes coupled to a voltage source for inducing a reaction to synthesize molecules in a respective test site.

35. The apparatus of Claim 1 wherein the array and electrodes are formed in integrated structures.

36. Apparatus for determining the presence of molecular structures within a substance comprising:

a) a source of said substance;

b) a plurality of sources of solutions containing known molecules which bind uniquely with different molecules; and

c) mixing means for selectively mixing each of said solutions with said substance; and

d) a detector for detecting the occurrence in the mixed solutions of binding between the known molecules and molecular structures in the substance.

37. The apparatus of Claim 36 wherein the detector detects binding by observing change in optical properties.

38. The apparatus of Claim 36 wherein the detector detects binding by observing change in electrical properties.

39. The apparatus of Claim 35 wherein the plurality of sources are contained in respective capillaries, each capillary having a valve for connecting a respective source to a stream of said substance.

40. The apparatus of Claim 39 wherein the capillaries and valves are formed in silicon and have diameters in the range of 1 to 10 microns.

41. Apparatus for synthesizing molecular structures comprising:

a) an array of test sites;

b) a source of chemical reactants disposed near the test sites;

c) electrodes associated with respective test sites; and

d) means for applying a potential to a respective electrode to attract said chemicals to a respective test site.

42. Apparatus for enhancing hybridization between a synthesized probe and a target molecule comprising:

a) an array of test sites each site containing a plurality of said probes;

b) an electrode associated with each site;

c) a source of target molecules applied to said sites; and

d) a voltage source for applying a potential to a respective electrode for attracting said target molecules to said probes.

43. The apparatus of Claim 11 wherein the test sites comprise wells formed in the substrate.

44. The apparatus of Claim 43 wherein the wells are formed with a textured surface.

45. The apparatus of Claim 44 wherein the textured surface consists of corrugations.

46. The apparatus of Claim 45 wherein the surface of the electrodes are also corrugated.

47. The apparatus of Claim 15 wherein the spacing between the electrode fingers in the lower portion and the electrode fingers in the upper portion is in the order of the diameter in solution of a target DNA molecule.

48. The method of Claim 6 wherein the dielectric property is permittivity.

49. The method of Claim 37 wherein the electrical property is permittivity.

50. The method of Claim 1 wherein the sample substance is in a solid state.

51. The method of Claim 8 wherein the resonant structure is a transmission line and changes in phase or amplitude of a signal propagated on the line are detected.

52. A method for determining the presence of molecular structures within a sample substance comprising:

a) forming an array of test sites having probes formed therein for binding to molecular structures, each site having probes which bind to a unique molecular structure;

b) dispensing a sample substance into a test site;

c) generating radiation through the test sites; and

d) detecting the difference in the radiation which is absorbed by the respective test sites to determine the presence of the molecular structures which are bound to a probe.

53. The method of Claim 52 wherein the difference is detected by an array of detectors formed of charge-coupled devices.

54. The method of Claim 53 wherein the array of detectors is formed integral with the array of test sites.

55. The method of Claim 53 wherein the array of detectors is formed separate from the array of test sites.

56. The method of Claim 55 wherein the array of detectors is aligned with the array of test sites and radiation is projected through the test sites onto the detector array.

57. The method of Claim 56 wherein the radiation is in the form of photons, or radioactive particles.

58. The method of Claim 52 wherein the radiation is generated within the test sites by a radioactive, chemical, thermal, chemiluminescent, or luminescent reaction.

59. The method of Claim 52 wherein the detectors detect thermal energy occurring when a bonding reaction takes place.

60. The apparatus of Claim 18 wherein the test sites include electrodes formed within wells formed in the plate.

61. The apparatus of Claim 60 wherein a surface of the well is textured.

62. The apparatus of Claim 61 wherein the texture is in the form of corrugations.

63. The apparatus of Claim 60 wherein a surface of an electrode is textured.

64. The apparatus of Claim 63 wherein the texture consists of corrugations.

65. A method for attaching probes to test sites formed in a substrate comprising the steps of:

a) forming test sites in a substrate;

b) forming adhesion material in said test sites to which said probes adhere; and

c) contacting said probes with said material.

66. The method of Claim 65 wherein a passivating layer covers the adhesion material and portions of the passivating layer are selectively removed to enable contact between the probe and the adhesive material at selected sites.

67. The method of Claim 66 wherein the portions are removed by laser ablation.

68. A method for attaching probes to a test site formed in a substrate comprising the steps of:

a) forming test sites in a substrate;

b) forming an adhesive material in the test site which material enables probes to attach to the test sites;

c) forming a protective coating over the adhesive material;

d) contacting the coating with a deprotecting material while selectively initiating a reaction at a selected site which removes the protective coating at the selected site; and

e) contacting the deprotected site with probes which adhere to the adhesive material.

69. The method of Claim 68 wherein the probes are presynthesized and the test sites consist of wells, the adhesive material is an epoxide, the protective coating is formed by hydrolyzing the epoxide, the deprotecting material is an acetate alcohol solution and the reaction is initiated at a selected cite by heating the site.

70. The method of Claim 68 wherein the reaction is initiated by selectively energizing resistors formed adjacent the test sites to heat the selected test sites.

71. The method of Claim 70 wherein non-selected test sites are maintained at a temperature above the desired reaction temperature.

72. The method of Claim 68 wherein the reaction is initiated by irradiating selected sites with light.

73. The method of Claim 72 wherein the light is in the visible or ultraviolet wavelength range and a photochemical reaction occurs.

74. The method of Claim 72 wherein the light originates from a laser which is scanned from site to site to initiate the reaction..

75. The method of Claim 72 wherein the light induces local heating of the site to initiate a reaction.

76. The method of Claim 72 wherein the light is generated by a light valve which projects light onto selected sites.

77. The apparatus of Claim 27 wherein the light source is a light valve for projecting light onto selected sites.

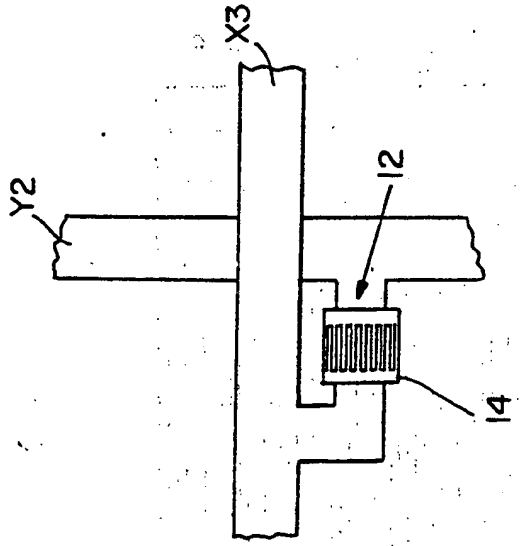
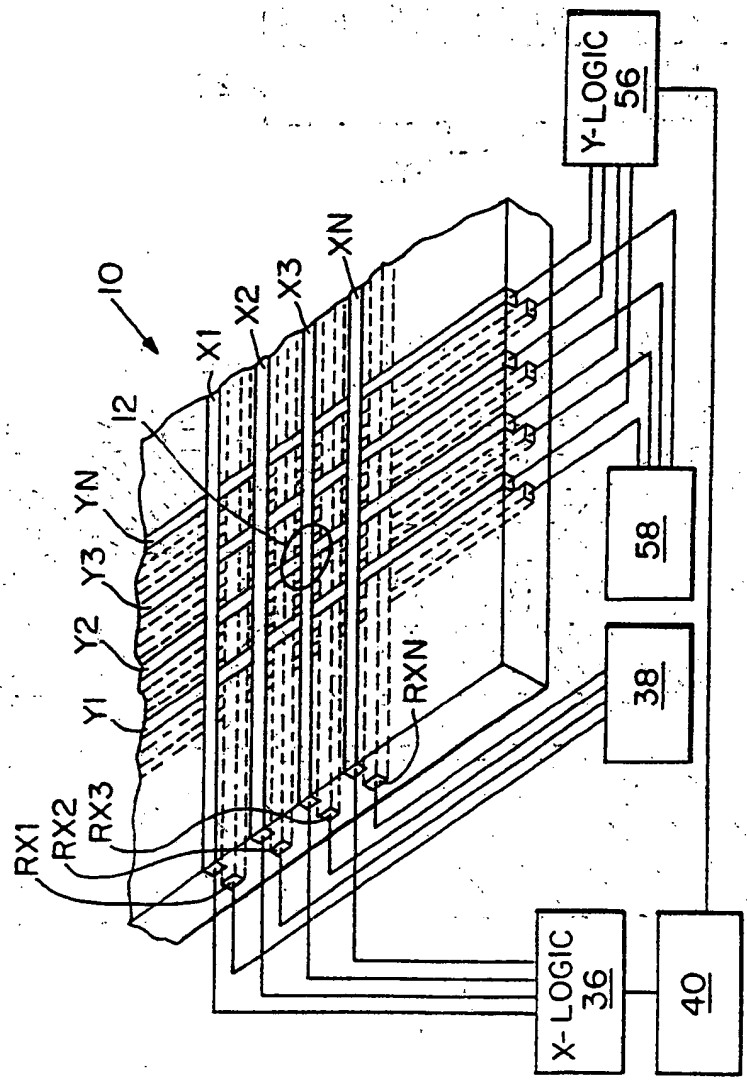

X3
12
Y2
14

Fig. 2


Y-LOGIC
56
58
38
X-LOGIC
36
40
10
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X2
X3
XN
12
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Y3
Y2
Y1
RXN
RX1
RX2
RX3

Fig. 1

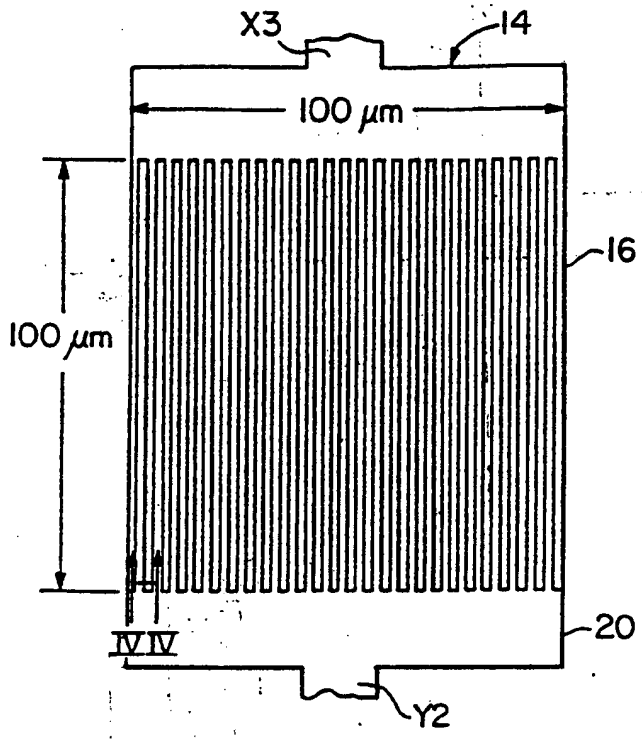

X3
14
100 μm
16
100 μm
20
IV IV
Y2

Fig. 3

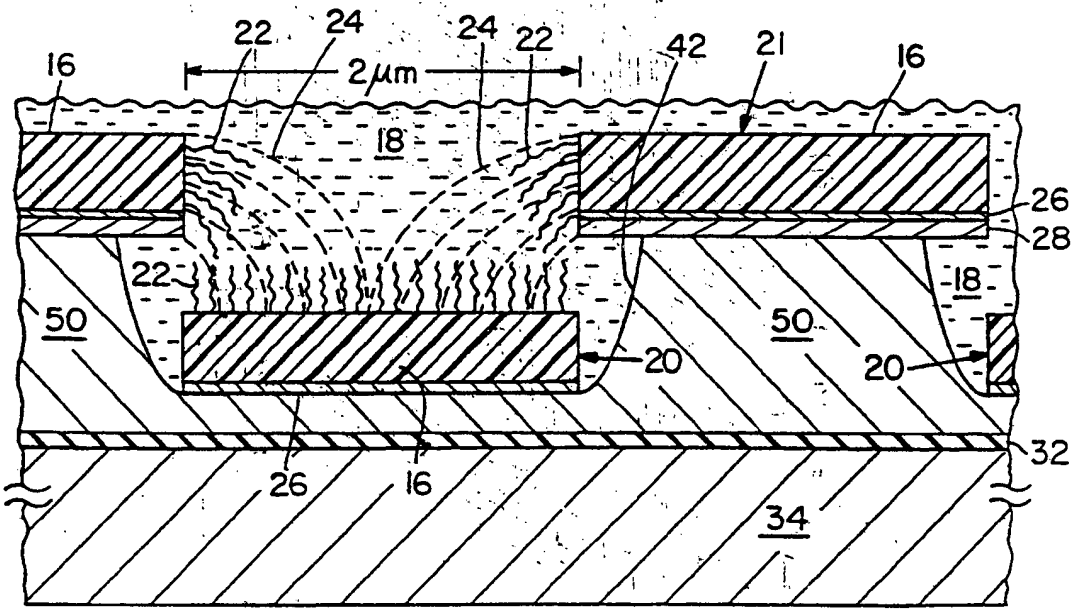

16
22
24
2 μm
24
22
42
21
16
18
26
28
22
18
50
50
20
20
32
26
16
34

Fig. 4

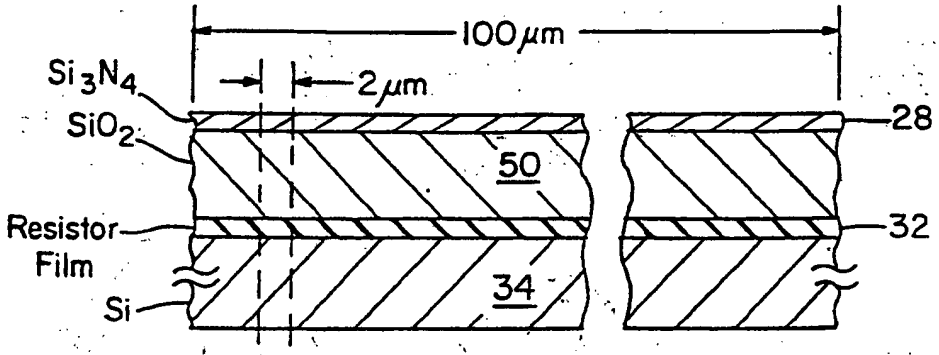


Fig. 5A

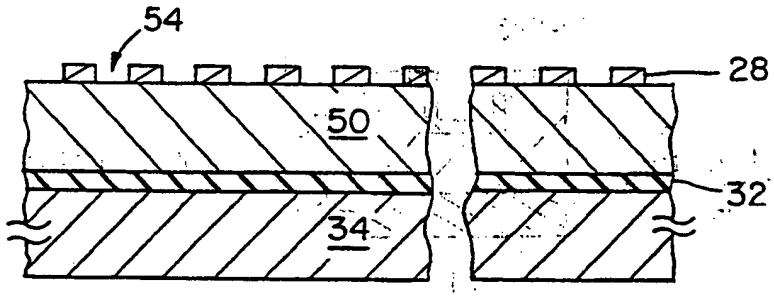


Fig. 5B

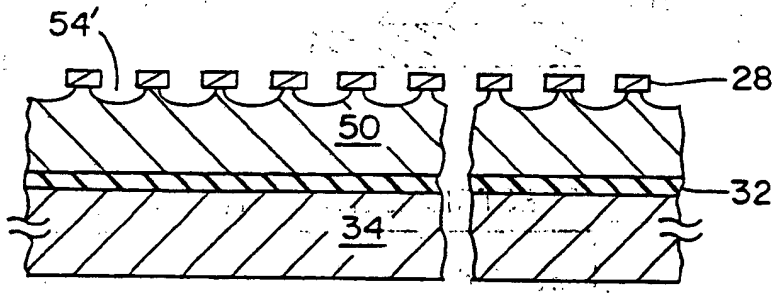


Fig. 5C

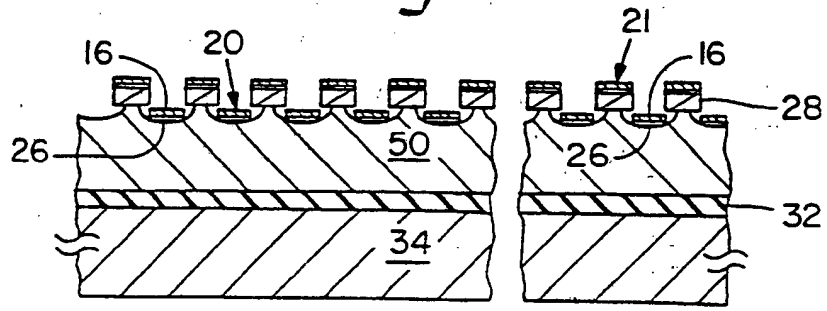


Fig. 5D

Fig. 6A

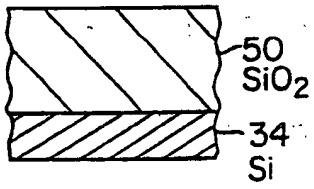


Fig. 6B

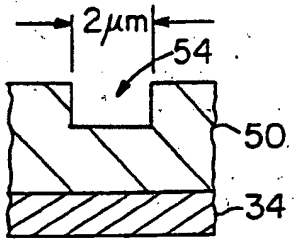


Fig. 6C

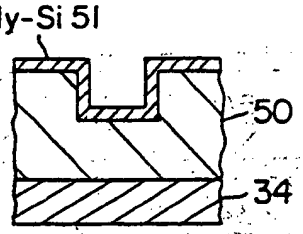


Fig. 6D

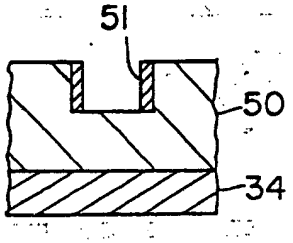


Fig. 6E

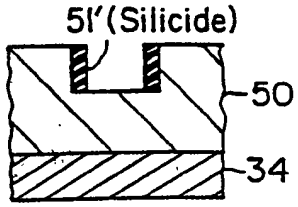


Fig. 6F

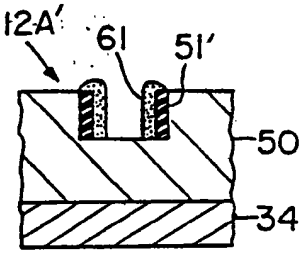


Fig. 6G

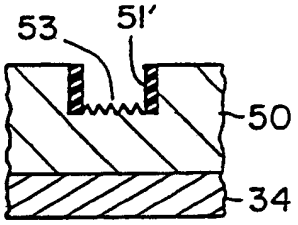
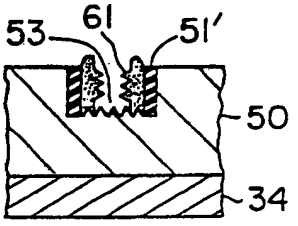


Fig. 6H



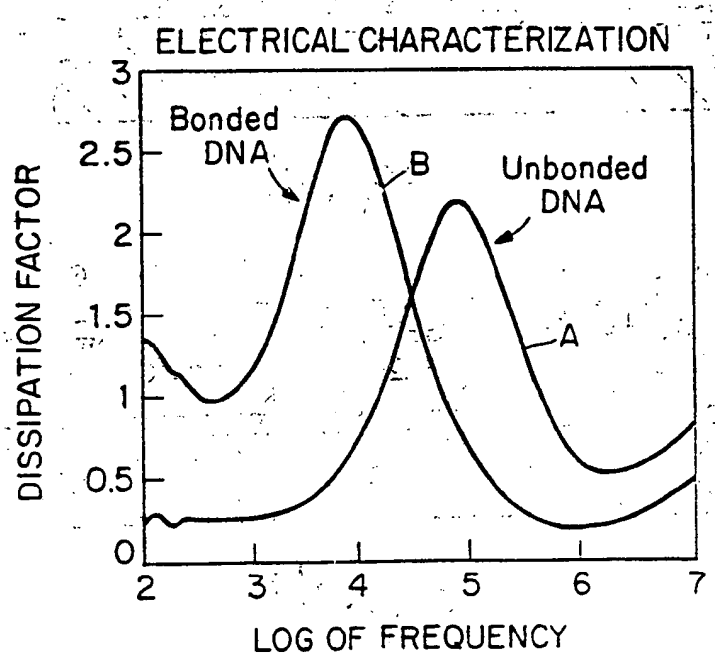


Fig. 7

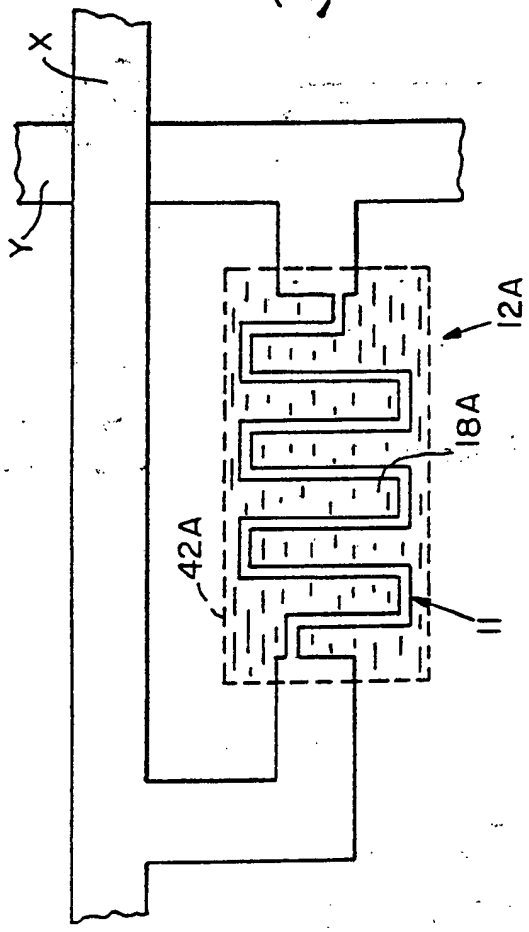


Fig. 8

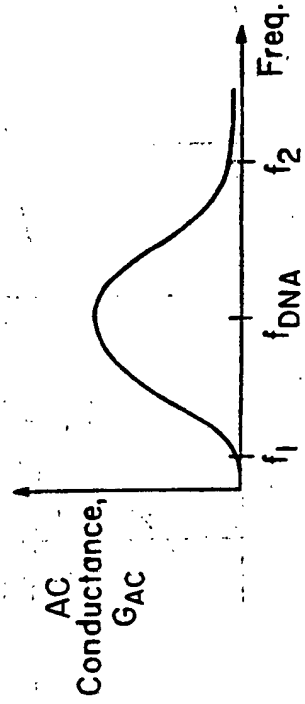


Fig. 10

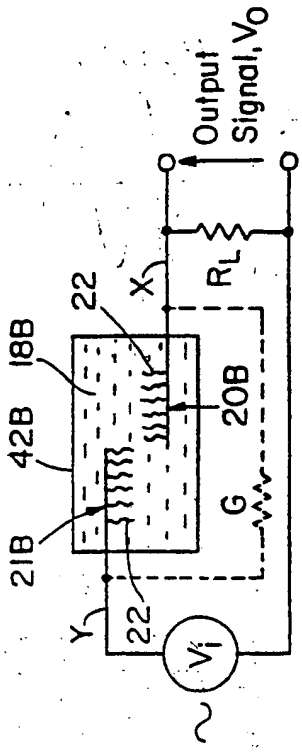


Fig. 9

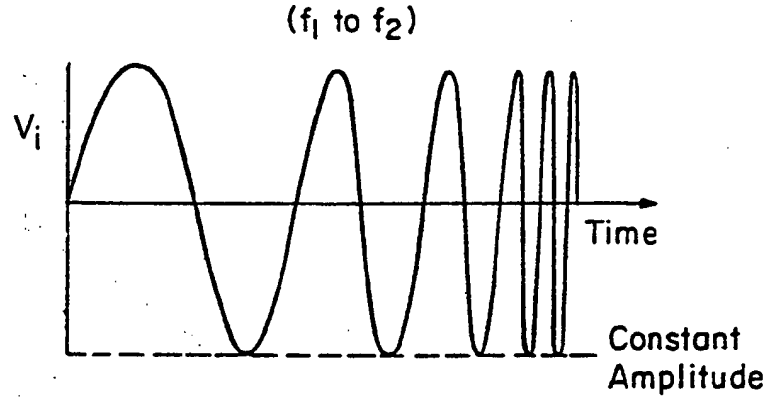


Fig. 11

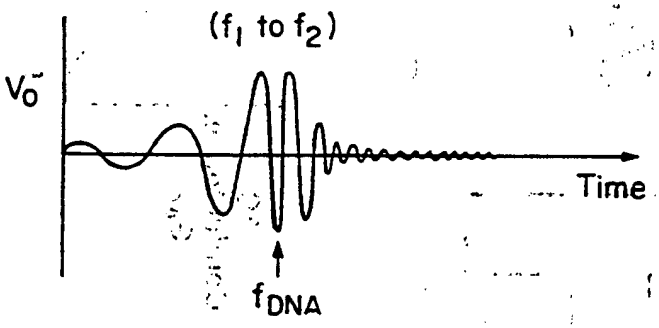


Fig. 12

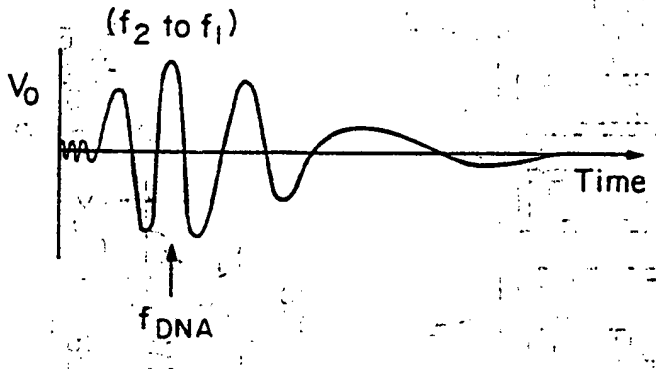


Fig. 13

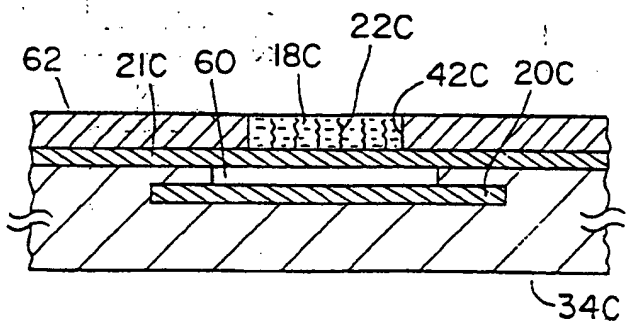


Fig. 14

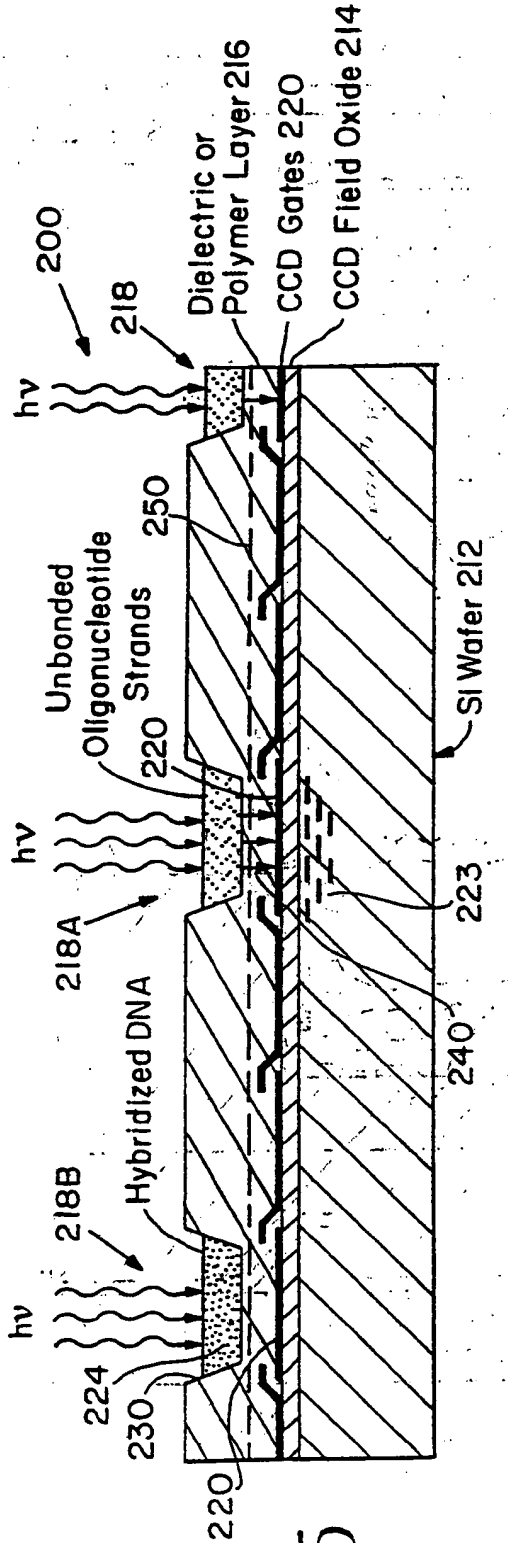

200
218
Dielectric or Polymer Layer 216
CCD Gates 220
CCD Field Oxide 214
hν
Unbonded Oligonucleotide Strands
250
220
Si Wafer 212
hν
218A
223
Hybridized DNA
240
218B
hν
224
230
220

Fig. 15

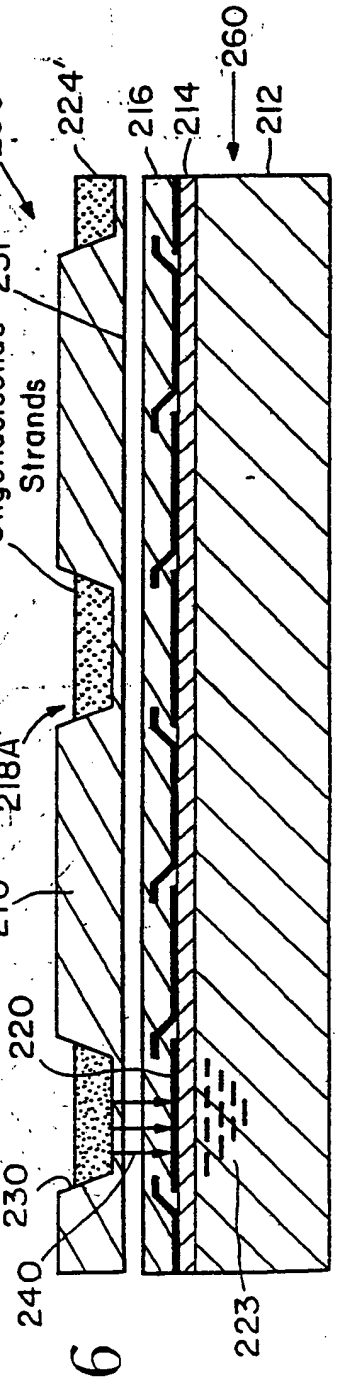

200'
224'
216
214
260
212
Unbonded Oligonucleotide Strands
251
218A'
270
220
230
240
223

Fig. 16

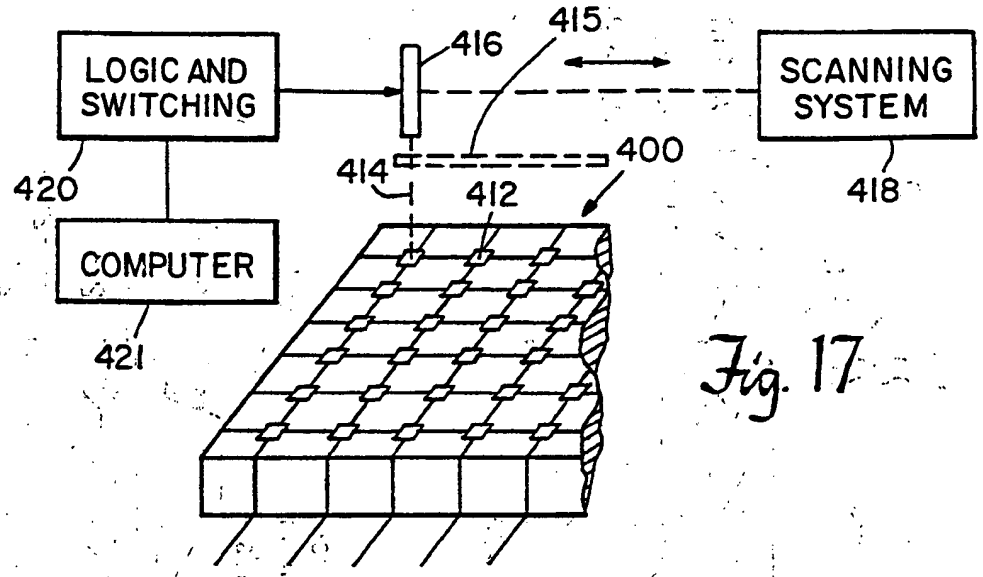

LOGIC AND SWITCHING
416
415
SCANNING SYSTEM
420
414
400
412
418
COMPUTER
421

Fig. 17

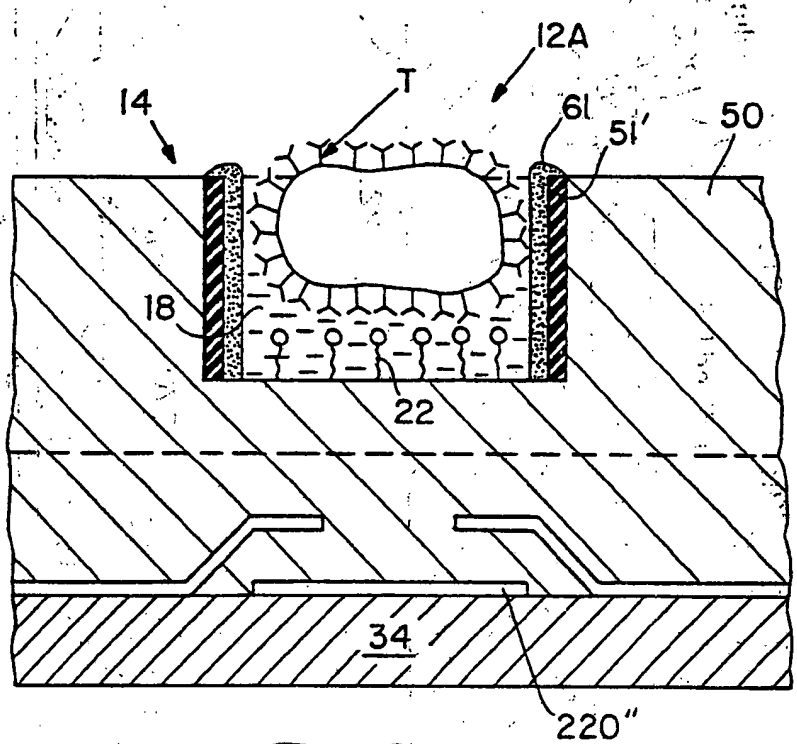

12A
T
14
61
51'
50
18
22
34
220"

Fig. 22

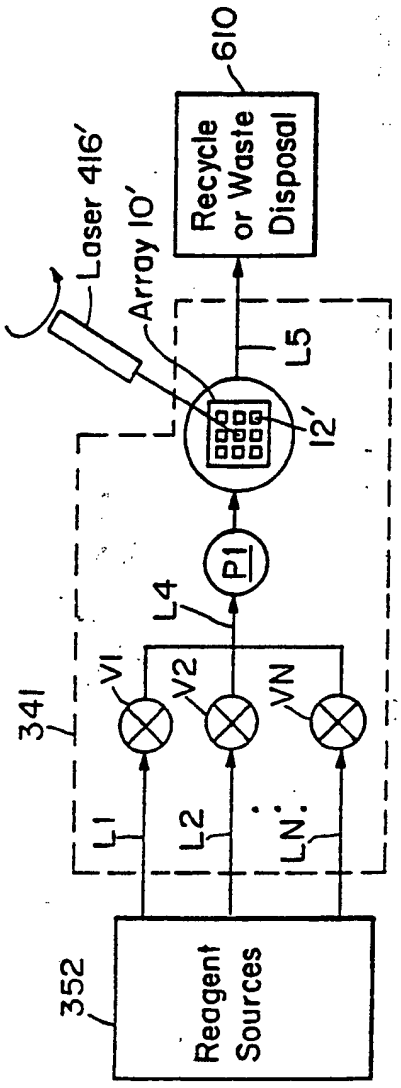

Laser 416'
Array 10'
Recycle or Waste Disposal
610
12'
L5
P1
L4
V1
V2
VN
341
L1
L2
LN
352
Reagent Sources

Fig. 18

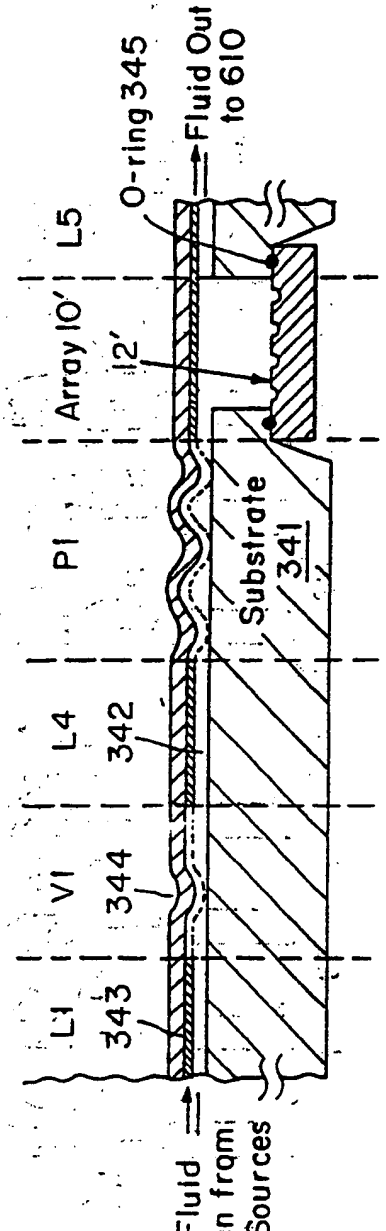

O-ring 345
Fluid Out to 610
L5
Array 10'
12'
P1
Substrate 341
L4
342
V1
344
L1
343
Fluid in from Sources

Fig. 19

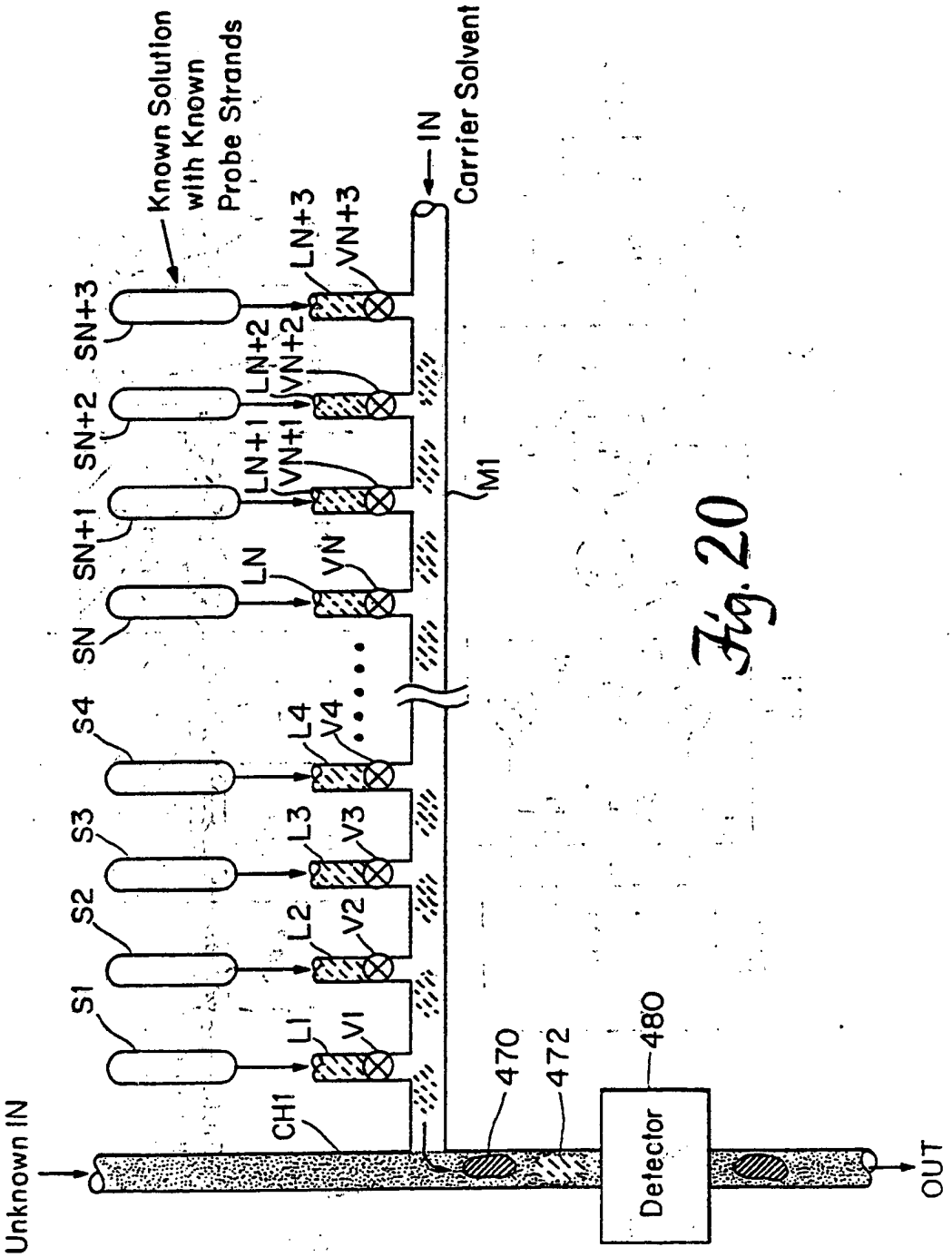

Known Solution with Known Probe Strands
IN
Carrier Solvent
SN+3
SN+2
SN+1
SN
S4
S3
S2
S1
LN+3
VN+3
LN+2
VN+2
LN+1
VN+1
LN
VN
L4
V4
L3
V3
L2
V2
L1
V1
M1
Unknown IN
CH1
470
472
480
Detector
OUT

Fig. 20

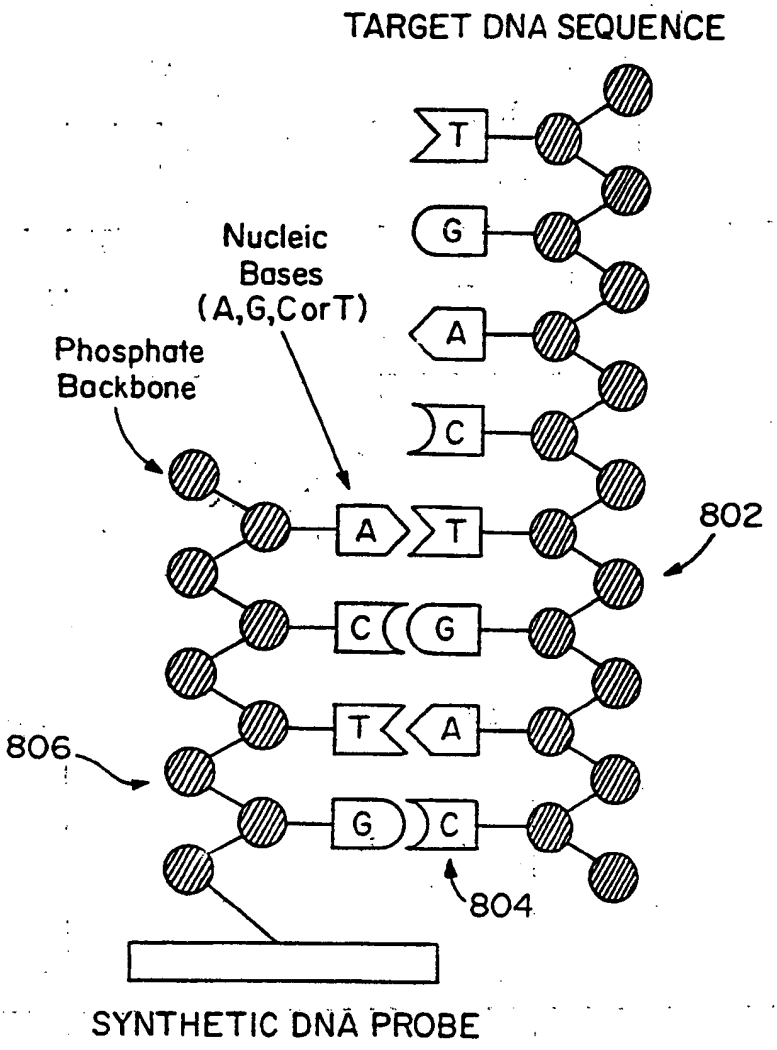

TARGET DNA SEQUENCE
T
G
A
C
Nucleic
Bases
(A,G,CorT)
Phosphate
Backbone
A
T
802
C
G
T
A
806
G
C
804
SYNTHETIC DNA PROBE

Fig. 21

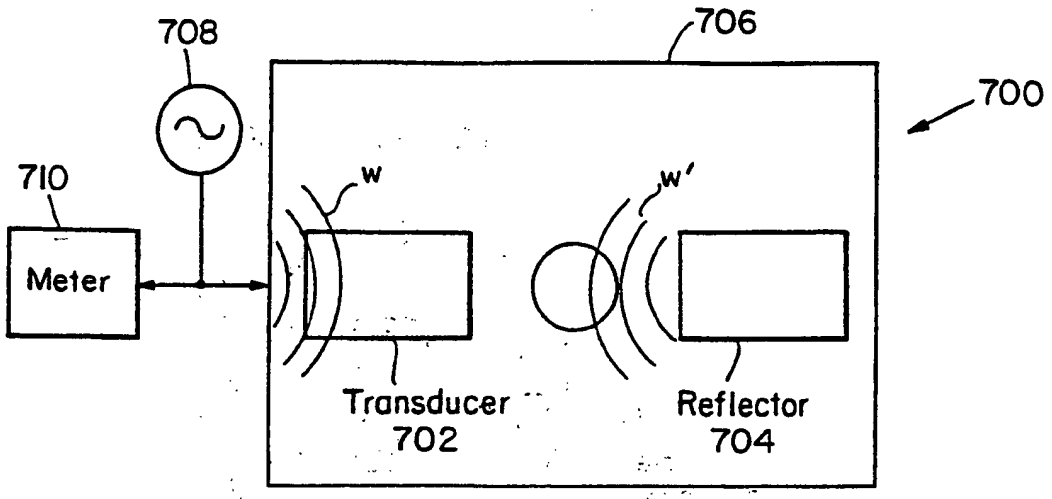

708
706
700
710
w
w'
Meter
Transducer
702
Reflector
704

Fig. 23

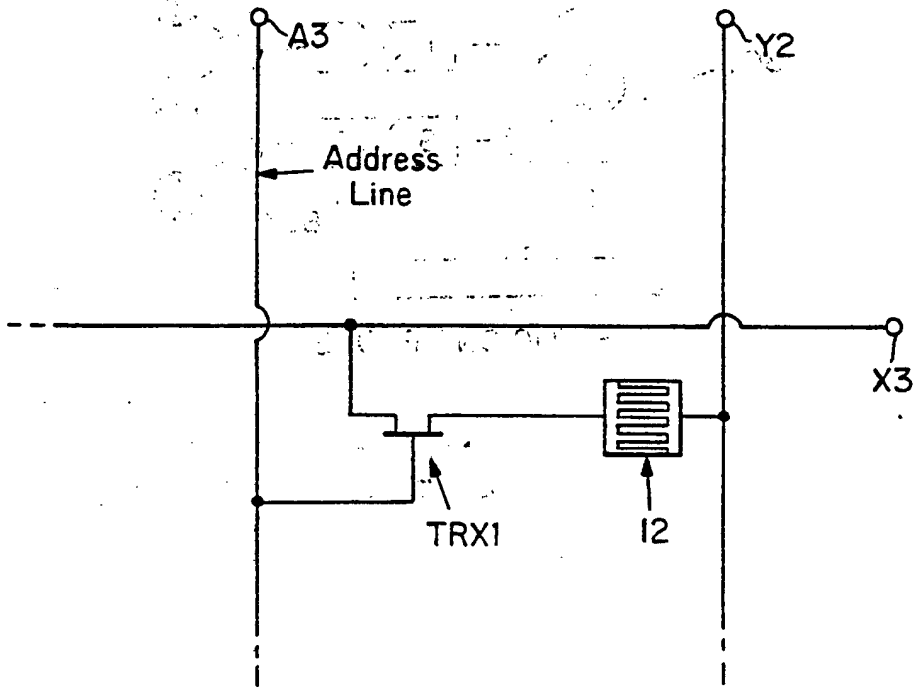

A3
Y2
Address
Line
X3
TRX1
12

Fig. 24

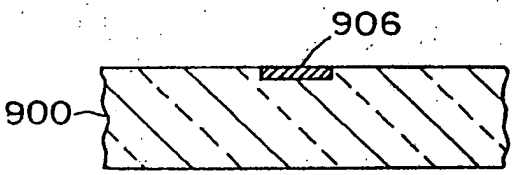


Fig. 25A

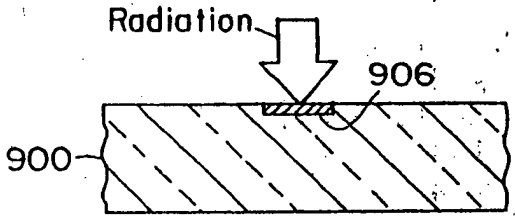


Fig. 25B

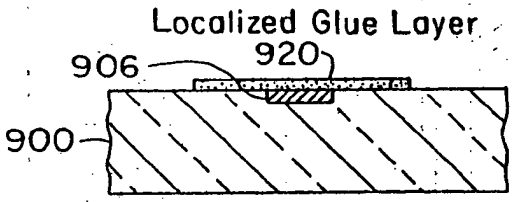


Fig. 25C

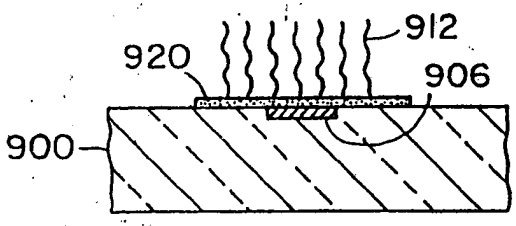


Fig. 25D

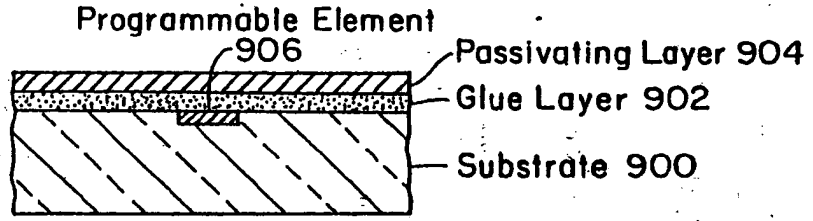


Fig. 26A

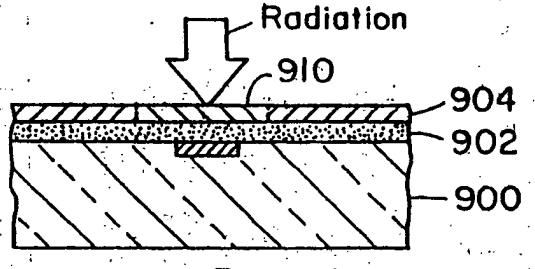


Fig. 26B

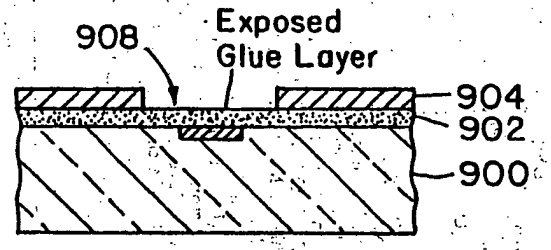


Fig. 26C

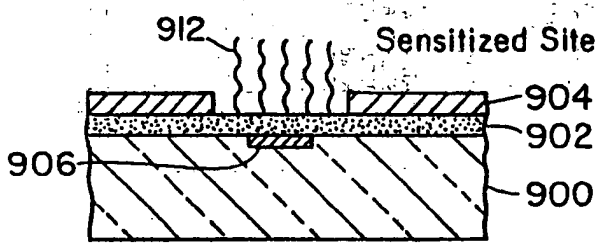


Fig. 26D

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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : G01N 33/543, C12Q 1/68 G01N 27/07, 21/75	A3	**(11) International Publication Number:** WO 93/22678 **(43) International Publication Date:** 11 November 1993 (11.11.93)

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07/872,582 23 April 1992 (23.04.92) US

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(74) Agents: REYNOLDS, Leo, R. et al.; Hamilton, Brook, Smith & Reynolds, Two Militia Drive, Lexington, MA 02173 (US).

(81) Designated States: CA, JP, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).

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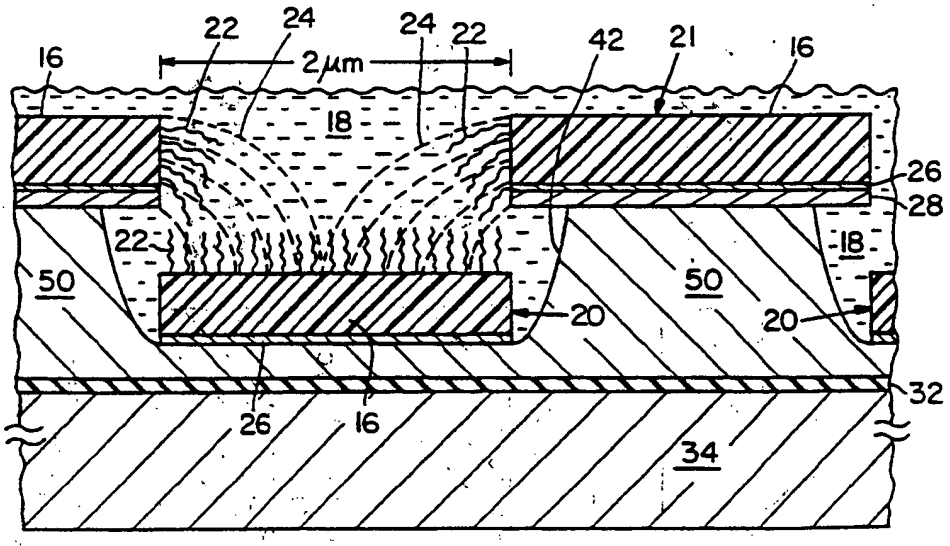
(54) Title: OPTICAL AND ELECTRICAL METHODS AND APPARATUS FOR MOLECULE DETECTION



(57) Abstract

A method and apparatus are disclosed for identifying molecular structures within a sample substance using a monolithic array of test sites formed on a substrate upon which the sample substance is applied. Each test site includes probes formed therein to bond with a predetermined target molecular structure or structures. A signal is applied to the test sites and certain electrical, mechanical and/or optical properties of the test sites are detected to determine which probes have bonded to an associated target molecular structure.

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INTERNATIONAL SEARCH REPORT

International Application No PCT/US 93/03829

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all)[6]

According to International Patent Classification (IPC) or to both National Classification and IPC

Int.Cl.5 G 01 N 33/543 C 12 Q 1/68 G 01 N 27/07
G 01 N 21/75

II. FIELDS SEARCHED

Minimum Documentation Searched[7]

Classification System	Classification Symbols
Int.Cl.5	B 01 L G 01 N C 12 Q

Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched[8]

III. DOCUMENTS CONSIDERED TO BE RELEVANT[9]

Category °	Citation of Document, [11] with indication, where appropriate, of the relevant passages [12]	Relevant to Claim No.[13]
X	WO,A,9015070 (AFFYMAX TECHNOLOGIES NV) 13 December 1990 cited in the application see page 20, line 6 - page 22, line 24 see page 28, line 14 - line 36 see page 39, line 1 - page 40, line 28; claim 22 ---	1,10,50
X	EP,A,0402917 (BIOCIRCUITS CORP) 19 December 1990	1,10,22
Y	see column 3, line 57 - column 4, line 27 see column 7, line 7 - line 57 see column 9, line 58 - column 11, line 16 see column 16, line 56 - column 21, line 25; figures 1,2 --- -/-	2

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IV. CERTIFICATION

Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report
14-09-1993	29. 12. 93
International Searching Authority	**Signature of Authorized Officer**
EUROPEAN PATENT OFFICE	BINDON C.A.

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)

Category °	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No.
Y	WO,A,9002327 (AUSTRALIAN MEMBRANE AND BIOTECHNOLOGY RESEARCH INSTITUTE) 8 March 1990	2
A	WO,A,9002327 see page 25, line 13 - page 27, line 35; figures 8-10	3-5
X	WO,A,9005300 (MIDWEST RESEARCH TECHNOLOGIES) 17 May 1990 see page 12, line 1 - page 18, line 8; figures 6-8	1,10,22
A	EP,A,0347579 (MESSERSCHMITT-B\LKOW-BLOHM GMBH) 27 December 1989 see column 2, line 41 - column 4, line 26 see column 7, line 2 - column 9, line 44	1,10,22
A	US,A,4963245 (WEETALL) 16 October 1990 see column 2, line 12 - column 3, line 13; figures	1-3,10, 22
P,A	US,A,5187096 (GIAEVER ET AL.) 16 February 1993 see column 2, line 33 - column 5, line 63; figures 1,2	1,2

INTERNATIONAL SEARCH REPORT

Int. ational application No.
PCT/US93/03829

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

For further information please see form PCT/ISA/206 mailed 27.09.93.

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2. ☐ As all searchable claims could be searches without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

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4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-5,10,22,50

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.

ANNEX TO THE INTERNATIONAL SEARCH REPORT ON INTERNATIONAL PATENT APPLICATION NO. US 9303829 SA 74062

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 16/12/93
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Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A- 9015070	13-12-90	US-A-	5143854	01-09-92
		AU-A-	5837190	07-01-91
		CA-A-	2054706	08-12-90
		EP-A-	0476014	25-03-92
		GB-A,B	2248840	22-04-92
		JP-T-	4505763	08-10-92
		NL-T-	9022056	02-03-92
EP-A- 0402917	19-12-90	US-A-	5156810	20-10-92
		CA-A-	2019039	15-12-90
		JP-A-	3128449	31-05-91
WO-A- 9002327	08-03-90	AU-A-	4078789	23-03-90
		EP-A-	0432188	19-06-91
		US-A-	5234566	10-08-93
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		CA-A-	2002660	10-05-90
EP-A- 0347579	27-12-89	DE-A-	3818614	07-12-89
		DE-A-	3825907	01-02-90
		US-A-	5252294	12-10-93
		DE-U-	8817007	02-10-91
US-A- 4963245	16-10-90	US-A-	5066372	19-11-91
US-A- 5187096	16-02-93	None		